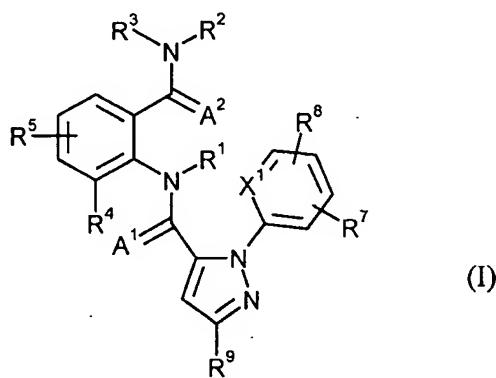


*Amendments to the Claims*

This listing of claims will replace all prior versions, and listings, of claims in the application.

1. (Currently amended) A composition comprising a synergistically effective active compound combination of at least one anthranilamide of the formula (I)



in which

A<sup>1</sup> and A<sup>2</sup> independently of one another represent oxygen or sulfur,

X<sup>1</sup> represents N or CR<sup>10</sup>,

R<sup>1</sup> represents hydrogen or represents C<sub>1</sub>-C<sub>6</sub>-alkyl, C<sub>2</sub>-C<sub>6</sub>-alkenyl, C<sub>2</sub>-C<sub>6</sub>-alkynyl or C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, each of which is optionally mono- or polysubstituted, where the substituents independently of one another may be selected from the group consisting of R<sup>6</sup>, halogen, cyano, nitro, hydroxyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, C<sub>1</sub>-C<sub>4</sub>-alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-alkylsulfonyl, C<sub>2</sub>-C<sub>4</sub>-alkoxycarbonyl, C<sub>1</sub>-C<sub>4</sub>-alkylamino, C<sub>2</sub>-C<sub>8</sub>-dialkylamino, C<sub>3</sub>-C<sub>6</sub>-cycloalkylamino, (C<sub>1</sub>-C<sub>4</sub>-alkyl)-C<sub>3</sub>-C<sub>6</sub>-cycloalkylamino and R<sup>11</sup>,

$R^2$  represents hydrogen,  $C_1$ - $C_6$ -alkyl,  $C_2$ - $C_6$ -alkenyl,  $C_2$ - $C_6$ -alkynyl,  $C_3$ - $C_6$ -cycloalkyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -alkylamino,  $C_2$ - $C_8$ -dialkylamino,  $C_3$ - $C_6$ -cycloalkylamino,  $C_2$ - $C_6$ -alkoxycarbonyl or  $C_2$ - $C_6$ -alkylcarbonyl,

$R^3$  represents hydrogen,  $R^{11}$  or represents  $C_1$ - $C_6$ -alkyl,  $C_2$ - $C_6$ -alkenyl,  $C_2$ - $C_6$ -alkynyl,  $C_3$ - $C_6$ -cycloalkyl, each of which is optionally mono- or polysubstituted, where the substituents independently of one another may be selected from the group consisting of  $R^6$ , halogen, cyano, nitro, hydroxyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy,  $C_1$ - $C_4$ -alkylthio,  $C_1$ - $C_4$ -alkylsulfinyl,  $C_1$ - $C_4$ -alkylsulfonyl,  $C_2$ - $C_6$ -alkoxycarbonyl,  $C_2$ - $C_6$ -alkylcarbonyl,  $C_3$ - $C_6$ -trialkylsilyl,  $R^{11}$ , phenyl, phenoxy and a 5- or 6-membered heteroaromatic ring, where each phenyl, phenoxy and 5- or 6-membered heteroaromatic ring may optionally be substituted and where the substituents independently of one another may be selected from one to three radicals  $W$  or one or more radicals  $R^{12}$ , or

$R^2$  and  $R^3$  may be attached to one another and form the ring  $M$ ,

$R^4$  represents hydrogen,  $C_1$ - $C_6$ -alkyl,  $C_2$ - $C_6$ -alkenyl,  $C_2$ - $C_6$ -alkynyl,  $C_3$ - $C_6$ -cycloalkyl,  $C_1$ - $C_6$ -haloalkyl,  $C_2$ - $C_6$ -haloalkenyl,  $C_2$ - $C_6$ -haloalkynyl,  $C_3$ - $C_6$ -halocycloalkyl, halogen, cyano, nitro, hydroxyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy,  $C_1$ - $C_4$ -alkylthio,  $C_1$ - $C_4$ -alkylsulfinyl,  $C_1$ - $C_4$ -alkylsulfonyl,  $C_1$ - $C_4$ -haloalkylthio,  $C_1$ - $C_4$ -haloalkylsulfinyl,  $C_1$ - $C_4$ -haloalkylsulfonyl,  $C_1$ - $C_4$ -alkylamino,  $C_2$ - $C_8$ -dialkylamino,  $C_3$ - $C_6$ -cycloalkylamino,  $C_3$ - $C_6$ -trialkylsilyl or represents phenyl, benzyl or phenoxy, each of which is optionally mono- or polysubstituted, where the substituents independently of one another may be

selected from the group consisting of C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>2</sub>-C<sub>4</sub>-alkenyl, C<sub>2</sub>-C<sub>4</sub>-alkynyl, C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>2</sub>-C<sub>4</sub>-haloalkenyl, C<sub>2</sub>-C<sub>4</sub>-haloalkynyl, C<sub>3</sub>-C<sub>6</sub>-halocycloalkyl, halogen, cyano, nitro, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, C<sub>1</sub>-C<sub>4</sub>-alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub>-alkylamino, C<sub>2</sub>-C<sub>8</sub>-dialkylamino, C<sub>3</sub>-C<sub>6</sub>-cycloalkylamino, C<sub>3</sub>-C<sub>6</sub>-(alkyl)cycloalkylamino, C<sub>2</sub>-C<sub>4</sub>-alkylcarbonyl, C<sub>2</sub>-C<sub>6</sub>-alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub>-alkylaminocarbonyl, C<sub>3</sub>-C<sub>8</sub>-dialkylaminocarbonyl and C<sub>3</sub>-C<sub>6</sub>-trialkylsilyl,

R<sup>5</sup> and R<sup>8</sup> in each case independently of one another represent hydrogen, halogen or represent in each case optionally substituted C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, R<sup>12</sup>, G, J, -OJ, -OG, -S(O)<sub>p</sub>-J, -S(O)<sub>p</sub>-G, -S(O)<sub>p</sub>-phenyl, where the substituents independently of one another may be selected from one to three radicals W or from the group consisting of R<sup>12</sup>, C<sub>1</sub>-C<sub>10</sub>-alkyl, C<sub>2</sub>-C<sub>6</sub>-alkenyl, C<sub>2</sub>-C<sub>6</sub>-alkynyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy and C<sub>1</sub>-C<sub>4</sub>-alkylthio, where each substituent may be substituted by one or more substituents independently of one another selected from the group consisting of G, J, R<sup>6</sup>, halogen, cyano, nitro, amino, hydroxyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, C<sub>1</sub>-C<sub>4</sub>-alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub>-haloalkylthio, C<sub>1</sub>-C<sub>4</sub>-haloalkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-haloalkylsulfonyl, C<sub>1</sub>-C<sub>4</sub>-alkylamino, C<sub>2</sub>-C<sub>8</sub>-dialkylamino, C<sub>3</sub>-C<sub>6</sub>-trialkylsilyl, phenyl and phenoxy, where each phenyl or phenoxy ring may optionally be substituted and where the substituents independently of one another may be selected from one to three radicals W or one or more radicals R<sup>12</sup>,

G in each case independently of one another represents a 5- or 6-membered non-aromatic carbocyclic or heterocyclic ring which may optionally contain one or two ring members from the group consisting of C(=O), SO and S(=O)<sub>2</sub> and which may optionally be substituted by one to four substituents independently of one another selected from the group consisting of C<sub>1</sub>-C<sub>2</sub>-alkyl, halogen, cyano, nitro and C<sub>1</sub>-C<sub>2</sub>-alkoxy, or independently of one another represents C<sub>2</sub>-C<sub>6</sub>-alkenyl, C<sub>2</sub>-C<sub>6</sub>-alkynyl, C<sub>3</sub>-C<sub>7</sub>-cycloalkyl, (cyano)-C<sub>3</sub>-C<sub>7</sub>-cycloalkyl, (C<sub>1</sub>-C<sub>4</sub>-alkyl)-C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, (C<sub>3</sub>-C<sub>6</sub>-cycloalkyl)-C<sub>1</sub>-C<sub>4</sub>-alkyl, where each cycloalkyl, (alkyl)cycloalkyl and (cycloalkyl)alkyl may optionally be substituted by one or more halogen atoms,

J in each case independently of one another represents an optionally substituted 5- or 6-membered heteroaromatic ring, where the substituents independently of one another may be selected from one to three radicals W or one or more radicals R<sup>12</sup>,

R<sup>6</sup> independently of one another represents -C(=E<sup>1</sup>)R<sup>19</sup>, -LC(=E<sup>1</sup>)R<sup>19</sup>, -C(=E<sup>1</sup>)LR<sup>19</sup>, -LC(=E<sup>1</sup>)LR<sup>19</sup>, -OP(=Q)(O R<sup>19</sup>)<sub>2</sub>, -SO<sub>2</sub>LR<sup>18</sup> or -LSO<sub>2</sub>LR<sup>19</sup>, where each E<sup>1</sup> independently of one another represents O, S, N-R<sup>15</sup>, N-OR<sup>15</sup>, N-N(R<sup>15</sup>)<sub>2</sub>, N-S=O, N-CN or N-NO<sub>2</sub>,

R<sup>7</sup> represents hydrogen, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, halogen, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, C<sub>1</sub>-C<sub>4</sub>-alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub>-haloalkylthio, C<sub>1</sub>-C<sub>4</sub>-haloalkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-haloalkylsulfonyl,

R<sup>9</sup> represents C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkylsulfinyl or halogen,

R<sup>10</sup> represents hydrogen, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, halogen, cyano or C<sub>1</sub>-C<sub>4</sub>-haloalkoxy,

R<sup>11</sup> in each case independently of one another represents in each case optionally mono- to trisubstituted C<sub>1</sub>-C<sub>6</sub>-alkylthio, C<sub>1</sub>-C<sub>6</sub>-alkylsulfenyl, C<sub>1</sub>-C<sub>6</sub>-haloalkylthio, C<sub>1</sub>-C<sub>6</sub>-haloalkylsulfenyl, phenylthio or phenylsulfenyl, where the substituents independently of one another may be selected from the list W, -S(O)<sub>n</sub>N(R<sup>16</sup>)<sub>2</sub>, -C(=O)R<sup>13</sup>, -L(C=O)R<sup>14</sup>, -S(C=O)LR<sup>14</sup>, -C(=O)LR<sup>13</sup>, -S(O)<sub>n</sub>NR<sup>13</sup>C(=O)R<sup>13</sup>, -S(O)<sub>n</sub>NR<sup>13</sup>C(=O)LR<sup>14</sup> or -S(O)<sub>n</sub>NR<sup>13</sup>S(O)<sub>2</sub>LR<sup>14</sup>,

L in each case independently of one another represents O, NR<sup>18</sup> or S,

R<sup>12</sup> in each case independently of one another represents -B(OR<sup>17</sup>)<sub>2</sub>, amino, SH, thiocyanato, C<sub>3</sub>-C<sub>8</sub>-trialkylsilyloxy, C<sub>1</sub>-C<sub>4</sub>-alkyl disulfides, -SF<sub>5</sub>, -C(=E<sup>1</sup>)R<sup>19</sup>, -LC(=E<sup>1</sup>)R<sup>19</sup>, -C(=E<sup>1</sup>)LR<sup>19</sup>, -LC(=E<sup>1</sup>)LR<sup>19</sup>, -OP(=Q)(OR<sup>19</sup>)<sub>2</sub>, -SO<sub>2</sub>LR<sup>19</sup> or -LSO<sub>2</sub>LR<sup>19</sup>,

Q represents O or S,

R<sup>13</sup> in each case independently of one another represents hydrogen or represents in each case optionally mono- or polysubstituted C<sub>1</sub>-C<sub>6</sub>-alkyl, C<sub>2</sub>-C<sub>6</sub>-alkenyl, C<sub>2</sub>-C<sub>6</sub>-alkynyl or C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, where the substituents independently of one another may be selected from the group consisting of R<sup>6</sup>, halogen, cyano, nitro, hydroxyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub>-

alkylamino, C<sub>2</sub>-C<sub>8</sub>-dialkylamino, C<sub>3</sub>-C<sub>6</sub>-cycloalkylamino or (C<sub>1</sub>-C<sub>4</sub>-alkyl)-C<sub>3</sub>-C<sub>6</sub>-cycloalkylamino,

R<sup>14</sup> in each case independently of one another represents in each case optionally mono- or polysubstituted C<sub>1</sub>-C<sub>20</sub>-alkyl, C<sub>2</sub>-C<sub>20</sub>-alkenyl, C<sub>2</sub>-C<sub>20</sub>-alkynyl or C<sub>3</sub>-C<sub>6</sub>-cycloalkyl, where the substituents independently of one another may be selected from the group consisting of R<sup>6</sup>, halogen, cyano, nitro, hydroxyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub>-alkylamino, C<sub>2</sub>-C<sub>8</sub>-dialkylamino, C<sub>3</sub>-C<sub>6</sub>-cycloalkylamino and (C<sub>1</sub>-C<sub>4</sub>-alkyl)-C<sub>3</sub>-C<sub>6</sub>-cycloalkylamino or represent optionally substituted phenyl, where the substituents independently of one another may be selected from one to three radicals W or one or more radicals R<sup>12</sup>,

R<sup>15</sup> in each case independently of one another represents hydrogen or represents in each case optionally mono- or polysubstituted C<sub>1</sub>-C<sub>6</sub>-haloalkyl or C<sub>1</sub>-C<sub>6</sub>-alkyl, where the substituents independently of one another may be selected from the group consisting of cyano, nitro, hydroxyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, C<sub>1</sub>-C<sub>4</sub>-alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub>-haloalkylthio, C<sub>1</sub>-C<sub>4</sub>-haloalkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-haloalkylsulfonyl, C<sub>1</sub>-C<sub>4</sub>-alkylamino, C<sub>2</sub>-C<sub>8</sub>-dialkylamino, C<sub>2</sub>-C<sub>6</sub>-alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub>-alkylcarbonyl, C<sub>3</sub>-C<sub>6</sub>-trialkylsilyl and optionally substituted phenyl, where the substituents independently of one another may be selected from one to three radicals W or one or more radicals R<sup>12</sup>, or N(R<sup>15</sup>)<sub>2</sub> represents a cycle which forms the ring M,

$R^{16}$  represents  $C_1$ - $C_{12}$ -alkyl or  $C_1$ - $C_{12}$ -haloalkyl, or  $N(R^{16})_2$  represents a cycle which forms the ring M,

$R^{17}$  in each case independently of one another represents hydrogen or  $C_1$ - $C_4$ -alkyl, or  $B(OR^{17})_2$  represents a ring in which the two oxygen atoms are attached via a chain having two to three carbon atoms which are optionally substituted by one or two substituents independently of one another selected from the group consisting of methyl and  $C_2$ - $C_6$ -alkoxycarbonyl,

$R^{18}$  in each case independently of one another represents hydrogen,  $C_1$ - $C_6$ -alkyl or  $C_1$ - $C_6$ -haloalkyl, or  $N(R^{13})(R^{18})$  represents a cycle which forms the ring M,

$R^{19}$  in each case independently of one another represents hydrogen or represents in each case mono- or polysubstituted  $C_1$ - $C_6$ -alkyl, where the substituents independently of one another may be selected from the group consisting of cyano, nitro, hydroxyl,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy,  $C_1$ - $C_4$ -alkylthio,  $C_1$ - $C_4$ -alkylsulfinyl,  $C_1$ - $C_4$ -alkylsulfonyl,  $C_1$ - $C_4$ -haloalkylthio,  $C_1$ - $C_4$ -haloalkylsulfinyl,  $C_1$ - $C_4$ -haloalkylsulfonyl,  $C_1$ - $C_4$ -alkylamino,  $C_2$ - $C_8$ -dialkylamino,  $CO_2H$ ,  $C_2$ - $C_6$ -alkoxycarbonyl,  $C_2$ - $C_6$ -alkylcarbonyl,  $C_3$ - $C_6$ -trialkylsilyl and optionally substituted phenyl, where the substituents independently of one another may be selected from one to three radicals W,  $C_1$ - $C_6$ -haloalkyl,  $C_3$ - $C_6$ -cycloalkyl or phenyl or pyridyl, each of which is optionally mono- to trisubstituted by W,

M in each case represents an optionally mono- to tetrasubstituted ring which, in addition to the nitrogen atom attached to the substituent pair  $R^{13}$  and  $R^{18}$ ,  $(R^{15})_2$

or  $(R^{16})_2$ , contains two to six carbon atoms and optionally additionally a further nitrogen, sulfur or oxygen atom, where the substituents independently of one another may be selected from the group consisting of  $C_1$ - $C_2$ -alkyl, halogen, cyano, nitro and  $C_1$ - $C_2$ -alkoxy,

W in each case independently of one another represents  $C_1$ - $C_4$ -alkyl,  $C_2$ - $C_4$ -alkenyl,  $C_2$ - $C_4$ -alkynyl,  $C_3$ - $C_6$ -cycloalkyl,  $C_1$ - $C_4$ -haloalkyl,  $C_2$ - $C_4$ -haloalkenyl,  $C_2$ - $C_4$ -haloalkynyl,  $C_3$ - $C_6$ -halocycloalkyl, halogen, cyano, nitro,  $C_1$ - $C_4$ -alkoxy,  $C_1$ - $C_4$ -haloalkoxy,  $C_1$ - $C_4$ -alkylthio,  $C_1$ - $C_4$ -alkylsulfinyl,  $C_1$ - $C_4$ -alkylsulfonyl,  $C_1$ - $C_4$ -alkylamino,  $C_2$ - $C_8$ -dialkylamino,  $C_3$ - $C_6$ -cycloalkylamino,  $(C_1$ - $C_4$ -alkyl)- $C_3$ - $C_6$ -cycloalkylamino,  $C_2$ - $C_4$ -alkylcarbonyl,  $C_2$ - $C_6$ -alkoxycarbonyl,  $CO_2H$ ,  $C_2$ - $C_6$ -alkylaminocarbonyl,  $C_3$ - $C_8$ -dialkylaminocarbonyl or  $C_3$ - $C_6$ -trialkylsilyl,

n in each case independently of one another represents 0 or 1,

p in each case independently of one another represents 0, 1 or 2,

where, if (a)  $R^5$  represents hydrogen,  $C_1$ - $C_6$ -alkyl,  $C_1$ - $C_6$ -haloalkyl,  $C_2$ - $C_6$ -haloalkenyl,  $C_2$ - $C_6$ -haloalkynyl,  $C_1$ - $C_4$ -haloalkoxy,  $C_1$ - $C_4$ -haloalkylthio or halogen and (b)  $R^8$  represents hydrogen,  $C_1$ - $C_6$ -alkyl,  $C_1$ - $C_6$ -haloalkyl,  $C_2$ - $C_6$ -haloalkenyl,  $C_2$ - $C_6$ -haloalkynyl,  $C_1$ - $C_4$ -haloalkoxy,  $C_1$ - $C_4$ -haloalkylthio, halogen,  $C_2$ - $C_4$ -alkylcarbonyl,  $C_2$ - $C_6$ -alkoxycarbonyl,  $C_2$ - $C_6$ -alkylaminocarbonyl or  $C_3$ - $C_8$  dialkylaminocarbonyl, (c) at least one substituent selected from the group consisting of  $R^6$ ,  $R^{11}$  and  $R^{12}$  if present and (d) if  $R^{12}$  is not present, at least one of the radicals  $R^6$  and  $R^{11}$  is different

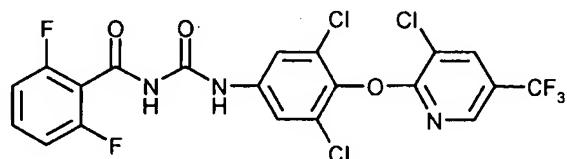
from C<sub>2</sub>-C<sub>6</sub>-alkylcarbonyl, C<sub>2</sub>-C<sub>6</sub>-alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub>-alkylaminocarbonyl and C<sub>3</sub>-C<sub>8</sub>-dialkylaminocarbonyl, and

where the compound of the general formula (I) may also be an N-oxide or salt.

and at least one insecticidally active compound of group 2 below, selected from

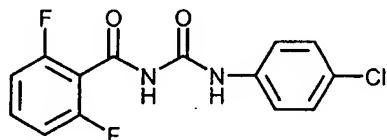
A) benzoylureas,

### (2-1) chlorfluazuron



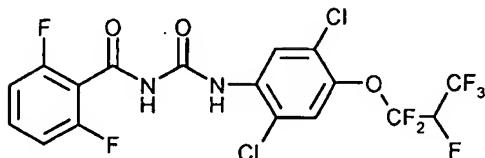
and/or

(2-2) diflubenzuron



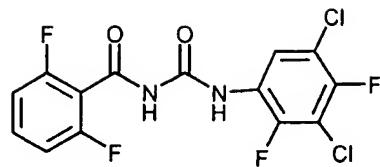
and/or.

(2-3) lufenuron



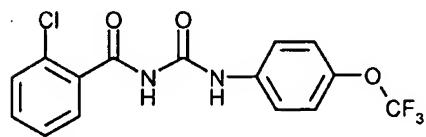
and/or

(2-4) teflubenzuron



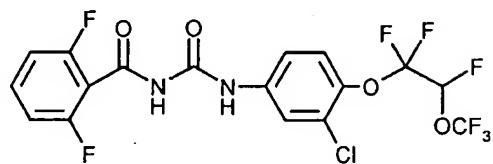
and/or

(2-5) triflumuron



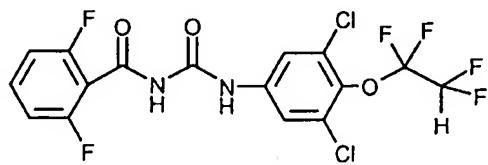
and/or

(2-6) novaluron



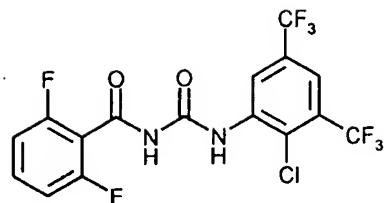
and/or

(2-7) hexaflumuron



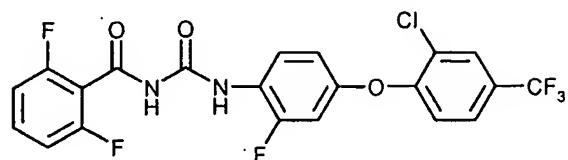
and/or

(2-8) bistrifluoron (DBI-3204)



and/or

(2-22) flufenoxuron



and/or

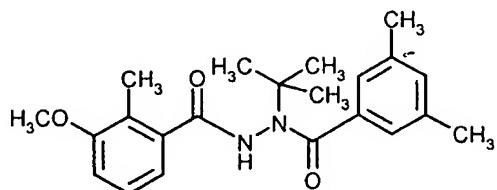
B) macrolides,

(2-9) emamectin

and/or

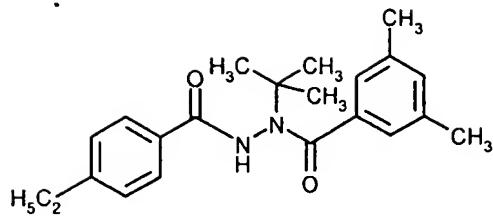
C) diacylhydrazines,

(2-10) methoxyfenozide



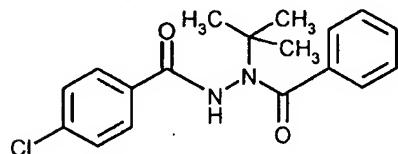
and/or

(2-11) tebufenozide



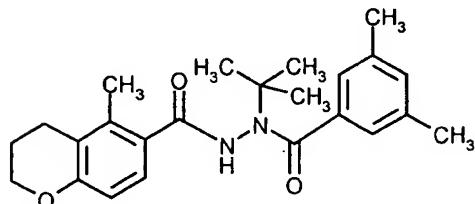
and/or

(2-12) halofenozide



and/or

(2-13) chromafenozide (ANS-118)



and/or

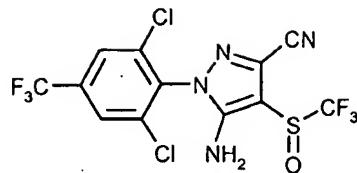
(2-14) *Trichogramma* spp.

and/or

(2-15) *Verticillium lecanii*

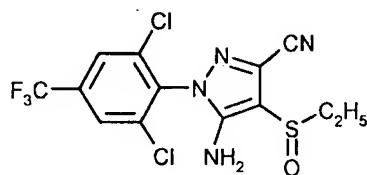
and/or

(2-16) fipronil



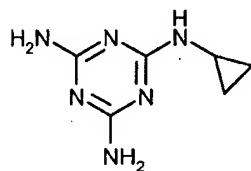
and/or

(2-17) ethiprole



and/or

(2-18) cyromazine

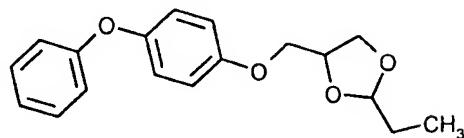


and/or

(2-19) azadirachtin

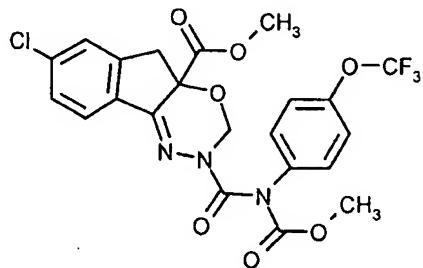
and/or

(2-20) diofenolan



and/or

(2-21) indoxacarb



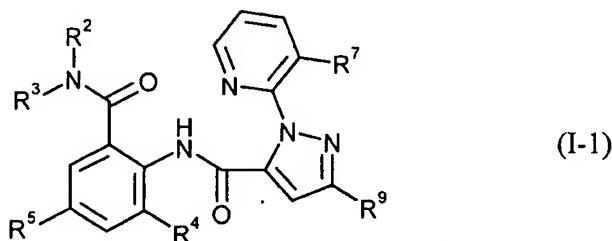
wherein the ratio of the at least one anthranilamide of formula (I) to the at least one active compound of group 2 is in a ratio of

<u>Formula (I) to chlorfluazuron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to diflubenzuron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to lufenuron</u>	<u>20 : 1 to 1 : 5</u>
<u>Formula (I) to teflubenzuron</u>	<u>20 : 1 to 1 : 5</u>
<u>Formula (I) to triflumuron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to novaluron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to hexaflumuron</u>	<u>20 : 1 to 1 : 5</u>
<u>Formula (I) to bistrifluoron</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to flufenoxuron</u>	<u>50 : 1 to 1 : 5</u>
<u>Formula (I) to emamectin</u>	<u>50 : 1 to 1 : 5</u>
<u>Formula (I) to methoxyfenozide</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to tebufenozide</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to halofenozide</u>	<u>2 : 1 to 1 : 100</u>
<u>Formula (I) to chromafenozide</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to Trichogramma spp.</u>	<u>1000 g a.i./ha : 20 000 wasps/ha</u> <u>to</u> <u>10 g a.i./ha : 500 000 wasps/ha</u>
<u>Formula (I) to Verticillium lecanii</u>	<u>0.05 % a.i. : 0.05 % F<sup>(*)</sup> to 0.001 %</u> <u>a.i. : 0.5 % F<sup>(*)</sup></u>
<u>Formula (I) to fipronil</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to ethiprole</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to cyromazine</u>	<u>10 : 1 to 1 : 10</u>
<u>Formula (I) to azadirachtin</u>	<u>50 : 1 to 1 : 5</u>

<u>Formula (I) to diofenolan</u>	<u>100 : 1 to 1 : 2</u>
<u>Formula (I) to indoxacarb</u>	<u>50 : 1 to 1 : 5</u>

F(\*) Formulation comprising 10<sup>9</sup> to 10<sup>10</sup> spores/g.

2. (Original) The composition as claimed in claim 1 comprising at least one active compound from the group of the anthranilamides of the formula (I-1) in which



in which

R<sup>2</sup> represents hydrogen or C<sub>1</sub>-C<sub>6</sub>-alkyl,  
R<sup>3</sup> represents C<sub>1</sub>-C<sub>6</sub>-alkyl which is optionally substituted by one R<sup>6</sup>,  
R<sup>4</sup> represents C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>2</sub>-haloalkyl, C<sub>1</sub>-C<sub>2</sub>-haloalkoxy or halogen,  
R<sup>5</sup> represents hydrogen, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>2</sub>-haloalkyl, C<sub>1</sub>-C<sub>2</sub>-haloalkoxy or halogen,  
R<sup>6</sup> represents -C(=E<sup>2</sup>)R<sup>19</sup>, -LC(=E<sup>2</sup>)R<sup>19</sup>, -C(=E<sup>2</sup>)LR<sup>19</sup> or -LC(=E<sup>2</sup>)LR<sup>19</sup>, where each E<sup>2</sup> independently of one another represents O, S, N-R<sup>15</sup>, N-OR<sup>15</sup>, N-N(R<sup>15</sup>)<sub>2</sub>, and each L independently of one another represents O or NR<sup>18</sup>,  
R<sup>7</sup> represents C<sub>1</sub>-C<sub>4</sub>-haloalkyl or halogen,  
R<sup>9</sup> represents C<sub>1</sub>-C<sub>2</sub>-haloalkyl, C<sub>1</sub>-C<sub>2</sub>-haloalkoxy, S(O)<sub>p</sub>-C<sub>1</sub>-C<sub>2</sub>-haloalkyl or halogen,

R<sup>15</sup> in each case independently of one another represents hydrogen or represents in each case optionally substituted C<sub>1</sub>-C<sub>6</sub>-haloalkyl or C<sub>1</sub>-C<sub>6</sub>-alkyl, where the substituents independently of one another may be selected from the group consisting of cyano, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, C<sub>1</sub>-C<sub>4</sub>-alkylthio, C<sub>1</sub>-C<sub>4</sub>-alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>-alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub>-haloalkylthio, C<sub>1</sub>-C<sub>4</sub>-haloalkylsulfinyl and C<sub>1</sub>-C<sub>4</sub>-haloalkylsulfonyl,

R<sup>18</sup> in each case represents hydrogen or C<sub>1</sub>-C<sub>4</sub>-alkyl,

R<sup>19</sup> in each case independently of one another represents hydrogen or C<sub>1</sub>-C<sub>6</sub>-alkyl,

p independently of one another represents 0, 1, 2.

3. (Previously presented) The composition as claimed in claim 1 wherein the at least one active compound of group 2 is selected from the group consisting of

(2-5) triflumuron

(2-22) flufenoxuron

(2-9) emamectin

(2-10) methoxyfenozide

(2-16) fipronil

(2-17) ethiprole and

(2-21) indoxacarb.

4. (Cancelled)

5. (Cancelled)
6. (Previously presented) A process for preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 1 is mixed with extenders and/or surfactants.
7. (Previously presented) A method for controlling animal pests, characterized in that synergistically effective mixtures as defined in claim 1 are allowed to act on animal pests and/or their habitat.
8. (Cancelled)
9. (Cancelled)
10. (Previously presented) A process for preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 2 is mixed with extenders and/or surfactants.
11. (Previously presented) A process for preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 3 is mixed with extenders and/or surfactants.

12. (Cancelled)

13. (Cancelled)

14. (Cancelled)

15. (Previously presented) A method for controlling animal pests, characterized in that synergistically effective mixtures as defined in claim 2 are allowed to act on animal pests and/or their habitat.

16. (Previously presented) A method for controlling animal pests, characterized in that synergistically effective mixtures as defined in claim 3 are allowed to act on animal pests and/or their habitat.

17. (Cancelled)

18. (Cancelled)

19. (Cancelled)

20. (Previously presented) The composition according to claim 1 wherein:  
 $R^2$  represents hydrogen or methyl,  
 $R^3$  represents  $C_1$ - $C_4$ -alkyl,

$R^4$  represents methyl, trifluoromethyl, trifluoromethoxy, fluorine, chlorine, bromine or iodine,

$R^5$  represents hydrogen, fluorine, chlorine, bromine, iodine, trifluoromethyl or trifluoromethoxy,

$R^7$  represents chlorine or bromine, and

$R^9$  represents trifluoromethyl, chlorine, bromine, difluoromethoxy or trifluoroethoxy.

21. (Previously presented) The composition according to claim 20 wherein the at least one active compound of group 2 is fipronil (2-16).

22. (Cancelled)

23. (Currently amended) The composition according to claim [[22]] 21 wherein the ratio of the at least one anthranilamide of formula (I) and fipronil (2-16) is from 5:1 to 1:5.

24. (Previously presented) The composition according to claim 23 wherein the ratio of the at least one anthranilamide of formula (I) and fipronil (2-16) is 1:5.

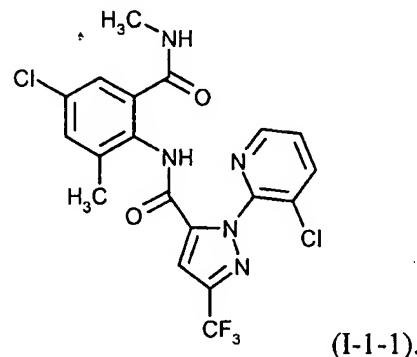
25. (Previously presented) The composition according to claim 20 wherein the at least one active compound of group 2 is ethiprole (2-17).

26. (Cancelled)

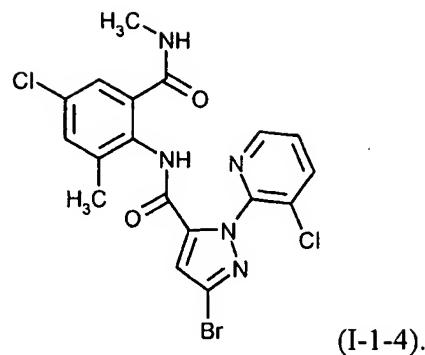
27. (Currently amended) The composition according to claim [[26]] 25 wherein the ratio of the at least one anthranilamide of formula (I) and ethiprole (2-17) is from 5:1 to 1:5.

28. (Previously presented) The composition according to claim 27 wherein the ratio of the at least one anthranilamide of formula (I) and ethiprole (2-17) is 1:5.

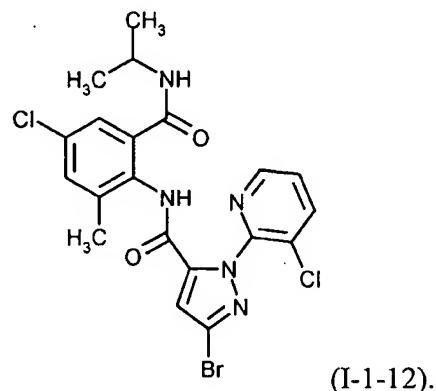
29. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



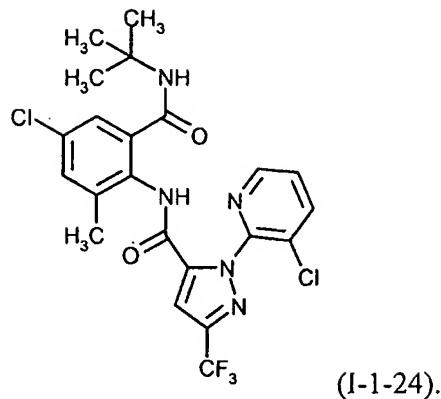
30. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



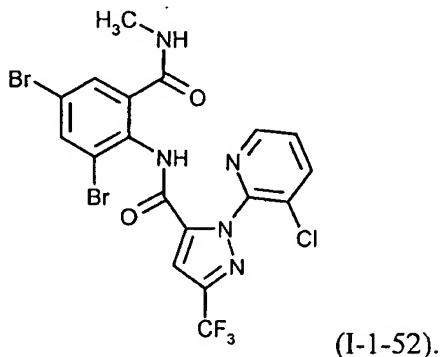
31. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



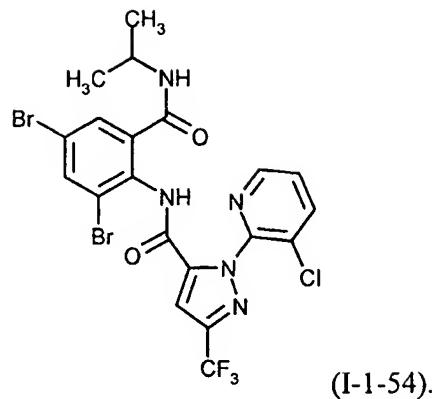
32. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



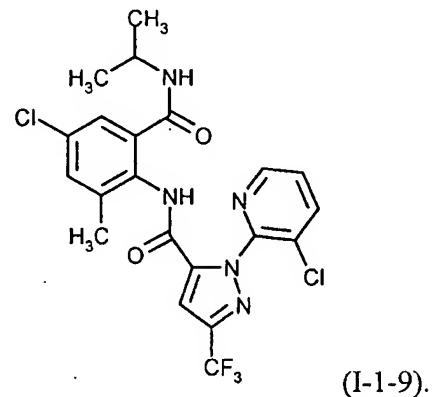
33. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



34. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



35. (Previously presented) The composition according to claim 20 wherein the at least one anthranilamide of formula (I-1) is



36. (Previously presented) The composition according to claim 35 wherein the at least one active compound of group 2 is fipronil (2-16).

37. (Previously presented) The composition according to claim 36 wherein the ratio of the compound (I-1-9) to fipronil (2-16) is 1:1.

38. (Previously presented) The composition according to claim 35 wherein the at least one active compound of group 2 is ethiprole (2-17).

39. (Previously presented) The composition according to claim 38 wherein the ratio of the compound (I-1-9) to ethiprole (2-17) is 1:1.

40. (New) The composition according to claim 1, wherein the ratio of the at least one anthranilamide of formula (I) to the at least one active compound of group 2 is in a ratio of

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Formula (I) to chlorfluazuron	5 : 1 to 1 : 5
Formula (I) to diflubenzuron	5 : 1 to 1 : 5
Formula (I) to lufenuron	10 : 1 to 1 : 2
Formula (I) to teflubenzuron	10 : 1 to 1 : 2
Formula (I) to triflumuron	5 : 1 to 1 : 5
Formula (I) to novaluron	5 : 1 to 1 : 5
Formula (I) to hexaflumuron	5 : 1 to 1 : 2
Formula (I) to bistrifluoron	5 : 1 to 1 : 5
Formula (I) to flufenoxuron	10 : 1 to 1 : 1
Formula (I) to emamectin	10 : 1 to 1 : 1
Formula (I) to methoxyfenozide	5 : 1 to 1 : 5
Formula (I) to tebufenozide	5 : 1 to 1 : 5
Formula (I) to halofenozide	1 : 1 to 1 : 30
Formula (I) to chromafenozide	5 : 1 to 1 : 5
Formula (I) to Trichogramma spp.	300 g a.i./ha : 50 000 wasps/ha to 50 g a.i./ha : 300 000 wasps/ha
Formula (I) to Verticillium lecanii	0.03 % a.i. : 0.1 % F <sup>(*)</sup> to 0.005 % a.i. : 0.2 % F <sup>(*)</sup>
Formula (I) to fipronil	5 : 1 to 1 : 5

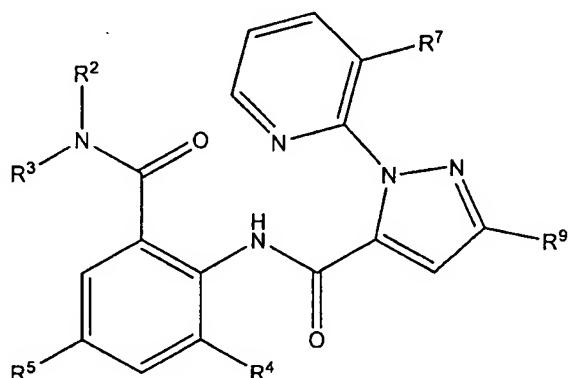
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Formula (I) to ethiprole	5 : 1 to 1 : 5
Formula (I) to cyromazine	5 : 1 to 1 : 5
Formula (I) to azadirachtin	10 : 1 to 1 : 1
Formula (I) to diofenolan	20 : 1 to 1 : 1
Formula (I) to indoxacarb	20 : 1 to 1 : 2

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F<sup>(\*)</sup> Formulation comprising 10<sup>9</sup> to 10<sup>10</sup> spores/g.

41. (New) A process of preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 40 is mixed with extenders and/or surfactants.
42. (New) A method for controlling animal pests, characterized in that a synergistically effective mixture as defined in claim 40 is allowed to act on animal pests and/or their habitat.
43. (New) The composition according to claim 1, wherein said compound of formula (I) is one of:



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Compound No.	R <sup>2</sup>	R <sup>3</sup>	R <sup>4</sup>	R <sup>5</sup>	R <sup>7</sup>	R <sup>9</sup>
I-1-1	H	Me	Me	Cl	Cl	CF <sub>3</sub>
I-1-4	H	Me	Me	Cl	Cl	Br
I-1-12	H	i-Pr	Me	Cl	Cl	Br
I-1-24	H	t-Bu	Me	Cl	Cl	CF <sub>3</sub>
I-1-52	H	Me	Br	Br	Cl	CF <sub>3</sub>
I-1-54	H	i-Pr	Br	Br	Cl	CF <sub>3</sub>

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44. (New) The composition according to claim 43, wherein said at least one compound of group 2 is ethiprole or fipronil.
45. (New) The composition according to claim 44, wherein the ratio of the compound of formula (1) to ethiprole or fipronil is from 5:1 to 1:5.
46. (New) A process of preparing pesticides, characterized in that a synergistically effective mixture as defined in claim 45 is mixed with extenders and/or surfactants.
47. (New) A method for controlling animal pests, characterized in that a synergistically effective mixture as defined in claim 45 is allowed to act on animal pests and/or their habitat.